

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
27 September 2001 (27.09.2001)

PCT

(10) International Publication Number
WO 01/70646 A1

(51) International Patent Classification⁷: C04B 28/04, 7/00

(21) International Application Number: PCT/EP01/03275

(22) International Filing Date: 21 March 2001 (21.03.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
00/03996 23 March 2000 (23.03.2000) FR

(71) Applicant (for all designated States except CA, FR, US):
SOFITECH N.V. [BE/BE]; Rue de Stalle 142, B-1180
Brussels (BE).

(71) Applicant (for CA only): SCHLUMBERGER CANADA
LIMITED [CA/CA]; Monenco Place, 24th floor, 801 6th
Avenue, S.W., Calgary, Alberta T2P 3W2 (CA).

(71) Applicant (for FR only): COMPAGNIE DES SER-
VICES DOWELL SCHLUMBERGER [FR/FR]; 50,
avenue Jean-Jaurès, F-92541 Montrouge (FR).

(72) Inventors; and

(75) Inventors/Applicants (for US only): BARLET-
GOUEDARD, Véronique [FR/FR]; Résidence La
Baleine, 2, impasse du Cachalot, F-92290 Chatenay Mal-
abry (FR). GOFFE, Bruno [FR/FR]; 1, rue Jeanne d'Arc,
F-91120 Palaiseau (FR).

(74) Agent: HYDEN, Martin; Etudes et Productions Schlum-
berger, Département Propriété Intellectuelle, 1, rue Henri
Becquerel, Boîte postale 202, F-92142 Clamart Cedex
(FR).

(81) Designated States (national): AL, AM, AT, AU, AZ, BA,
BB, BG, BR, BY, CA, CH, CN, CO, CU, CZ, DE, DK, EE,
ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD,
MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD,
SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ,
VN, YU, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: CEMENTING COMPOSITIONS AND APPLICATION OF THESE COMPOSITIONS TO CEMENTING OIL WELLS
OR THE LIKE

(57) Abstract: The present invention concerns cementing compositions for an oil well or the like, based on a Portland cement, silica and alumina, wherein the mineralogical composition of the cement matrix is included in the Si-Ca-Al triangle in one of the margarite-haüyne-[epidote/pumpellyite], haüyne-prehnite-[epidote/pumpellyite] and haüyne-prehnite-pectolite composition triangles. More particularly, the cementing compositions of the invention are suitable for cementing wells subjected to high temperatures and to chemical attack by brines, and are more suitable when the brine - or a saline formulation close to that of the brine - is used as the mixing water.

WO 01/70646 A1



CEMENTING COMPOSITIONS AND APPLICATION OF THESE COMPOSITIONS TO CEMENTING OIL WELLS OR THE LIKE.

5 The present invention relates to techniques for drilling oil, gas, water or geothermal wells or the like. More precisely, the invention relates to cementing compositions, more particularly suitable for high temperature wells such as geothermal wells, very deep wells or more generally, wells subjected to thermal and/or chemical attack, in particular as regards injecting steam to stimulate production or due to the presence of brine.

10

After drilling an oil well or the like, a casing or a coiled tubing is lowered into the hole and cemented over all or part of its depth. In particular, cementing prevents exchange of fluids between the different layers of formation traversed by the hole, prevents gas from rising in the annular space surrounding the casing or limits the ingress of water into the production well. Its principal purpose, of course, is to support the casing.

15

In the vast majority of cases, a Portland cement is used as the hydraulic binder, a material which is essentially constituted by calcium silicates and calcium aluminosilicates. In the presence of water, those mineral phases hydrate and preferentially form hydrated calcium silicate (CSH) and also portlandite ($\text{Ca}(\text{OH})_2$) and aluminium sulphate. CSH is a gel which is largely responsible for the cohesion and mechanical strength of the cement in the short to medium term.

20

25 When the temperature exceeds about 100°C , in the first days or months the majority of cements lose compressive strength and become permeable. Then they regain their strength and their permeability reduces. That phenomenon, known as retrogression, is due to metamorphosis of CSH gel to a highly crystalline phase known as alpha dicalcium silicate, $\alpha\text{-C}_2\text{SH}$, which is much more dense than CSH with a contraction in volume and an increase in the porosity of the cement. This results in disintegration of the cement which increases in rate as the porosity increases, rendering it more sensitive to chemical attack such as channelling of brine or circulation of carbon dioxide.

30

35 To avoid the phenomenon of retrogression, silica in the form of sand or silica flour is systematically added to compositions which have to withstand temperatures of more than 100°C . Adding 35% to 40% of silica (with

respect to the weight of the Portland cement) brings the CaO/SiO_2 ratio (usually denoted C/S) to about 1.

Under those conditions, CSH then transforms at about 110°C to tobermorite $\text{C}_3\text{S}_6\text{H}_5$, then to xonotlite $\text{C}_6\text{S}_6\text{H}$ if the temperature is raised to 150°C . Those phases ensure the stability of the cement at high temperature, in the absence of brine. Those two minerals are in themselves very strong but they have slightly different apparent densities, which destabilises the cement. Further, near 250°C , a new mineralogical phase, truscottite ($\text{C}_7\text{S}_{12}\text{H}_3$) forms which further embrittles the cement and increases its permeability. The degradation mechanisms described above are certainly not exhaustive, not least because the roles of other oxides present in the Portland cement, in particular alumina and ferrites, have been ignored, but they suffice to demonstrate the complexity of the phenomena occurring, without even considering the reaction products between the cement matrix and brines. In nature, in a geological medium, cement is thus metastable. In contact with brine, its mineralogical repertoire is modified by ion exchanges, which modify its physical qualities.

The problems of cement disintegration when the temperature is raised are more severe when developing very deep wells, or injecting steam to recover highly viscous oils, or in geothermal wells to exploit geothermal energy.

French patent FR-A-2 765 571 proposes selecting a mineralogical phase which is stable at the envisaged temperatures and pressures and promoting its formation by adding a certain quantity of a mineral oxide to a base constituted by Portland cement and silica. In particular, that patent describes compositions which can promote the formation of calcium hydrogarnets from the series $[\text{SiO}_4]\text{Ca}_3\text{M}^{2+}$, where $\text{M} = \text{Al}$ (grossularite) or $\text{M} = \text{Fe}$ (andradite) by adding alumina or ferrite. The quantity added is typically in the range 20% to 50% (by weight with respect to the weight of Portland cement); i.e., of the same order of magnitude as the amount of silica added. A more particularly preferred composition is constituted by class G Portland cement, 30% to 40% of silica (preferably of the order of 35%) and 20% to 40% of alumina (preferably of the order of 30%).

The mineral phases favoured by these compositions are principally of varying composition, in particular hydrogarnets from the hydrogrossularite series, with a structure derived from calcium garnets $\text{Ca}_3\text{Al}_2\text{Si}_3\text{O}_{12}$ (C_3AS_3) with the silica atoms omitted and replacement of the oxygen atoms to which they were bonded by hydroxyl

groups. When all of the silicon atoms are replaced, C_3AH_6 is obtained. Under the selected experimental conditions (288°C, 20 MPa), those garnets have unfortunately been shown to be poorly constituted in part, with low reproductivity of crystallisation leading to larger or smaller crystals associated with high heterogeneity in compression
5 test results.

The aim of the invention is to develop novel formulations suitable for cementing oil, gas, geothermal wells or the like, based on a Portland and alumina type cement tending to form mineral phases such that the properties of the cement matrix,
10 in particular regarding compressive strength, do not degrade in practice under the effect of a rise in temperature and pressure (250°C-300°C; 20 MPa) and/or chemical attack (brines).

The invention thus provides a cement matrix included in the Si-Ca-Al triangle
15 in one of the margarite-haüyne-[epidote/pumpellyite], haüyne-prehnite-[epidote/pumpellyite] and haüyne-prehnite-pectolite composition triangles. In the Si-Al-Ca-Na system, there exist in the triangles defined many mineral phases which are stable under the temperatures and pressures under consideration.

Preferably again, the compositions of the invention are located in the most
20 calcium-rich zones, in particular close to the margarite-[epidote/pumpellyite]-pectolite junctions and margarite-prehnite-pectolite junctions, corresponding to compositions within the Si-Ca-Al triangle approximately in the zone delimited by silica between 35% and 50%, aluminium between 20% and 38% and calcium between 25% and
25 35%.

Such mineral compositions are obtained, for example, by adding, to a Portland cement, a silica (typically 75% to 85% by weight with respect to the weight of class G Portland cement) and alumina (in the range 30% to 80% by weight with respect to the
30 weight of class G Portland cement). Preferably, all or part of the added silica is in the form of micro-silica, i.e., in a highly reactive form. The presence of a high reactivity phase in the formulation appears to encourage aluminosilicate crystallisation reactions during the first hours of setting, subsequently limiting competition between the kinetics of the dissolution of low or poor crystallinity initial phases and the kinetics of
35 aluminosilicate crystallisation.

The degradation kinetics in brine of the compositions of the invention are much slower than that of ordinary oil-well cements. Further, the presence of a brine during setting, or even deliberately introduced into the mixing water, is not a factor in rapid cement degradation. In contrast, the presence of sulphate on setting of the cement can encourage the formation of calco-sodium crystals which are particularly stable under the envisaged conditions.

Further advantageous details and characteristics of the invention will become apparent from the description below of tests established for different examples of additive compositions, made with reference to the figures which show:

Figure 1: a phase diagram of a ternary silica/aluminium/calcium system;

Figure 2: a graph showing the evolution of compressive strength after high temperature treatment in the presence of brine for different cement compositions.

The studies reported in French patent application FR-A-2 765 271 tended to promote mineral phases such as hydrogarnets from the hydrogrossularite series, C_3AH_6 , by modifying the composition of a hydraulic binder based on a Portland cement by adding a certain quantity of alumina. However, while various garnets are effectively formed under the selected experimental conditions (288°C, 20 MPa), it has been observed that these garnets were quite poorly constituted and of low durability and further, xonotlite persists in forming. Xonotlite embrittles cement by forming cavities inside the structure; further, this phenomenon is aggravated in the presence of brine by dissolution of xonotlite.

In order to better appreciate the adaptation possibilities of the system, Figure 1 shows the mineral phases of the Si-Al-Ca-Na system, summarised in Table I below and projected on the ternary Si-Al-Ca diagram. Clearly there exist many junctions between minerals which theoretically allow the xonotlite to disappear while avoiding the formation of garnets which have been proved to be unstable.

Classes	Structural forms	Classes	Structural forms
Plagioclase:	Albite $[\text{Si}_3\text{AlO}_8]\text{Na}$	Zeolite:	Analcime $[\text{Si}_2\text{AlO}_6](\text{Na}_2, \text{Ca})2\text{H}_2\text{O}$
	Anorthite $[\text{Si}_2\text{Al}_2\text{O}_8]\text{Ca}$		Gismondine $[\text{Si}_2\text{Al}_2\text{O}_8]\text{Ca}4\text{H}_2\text{O}$
Feldspathoid:	Cancrinite $[\text{SiAlO}_4]_3\text{CO}_3\text{Na}_3\text{Ca}$		Heulandite $[\text{Si}_7\text{Al}_2\text{O}_{18}]\text{Ca}6\text{H}_2\text{O}$
	Häüyne $[\text{SiAlO}_4]_6(\text{SO}_4, \text{S})_1$ $2(\text{Na}, \text{Ca})_{4.8}$		Laumontite $[\text{Si}_2\text{AlO}_6]\text{Ca}_4\text{H}_2\text{O}$
	Noselite $[\text{SiAlO}_4]_6\text{SO}_4\text{Na}_8$		Pectolite $[\text{SiO}_3]_3\text{NaCa}_2\text{H}$
	Sodalite $[\text{SiAlO}_4]_6\text{Cl}_2\text{Na}_8$		
Micas:	Margarite $[\text{SiAlO}_3]_2(\text{OH})_2\text{CaAl}_2$		Prehnite $[\text{Si}_3\text{AlO}_{10}]\text{Ca}_2\text{Al}(\text{OH})_2$
Epidote:	Zoisite $[\text{SiO}_3][\text{SiO}_4]$ $(\text{OH})\text{Al}_2\text{O CaAl}_2$		Wairakite $[\text{Si}_2\text{AlO}_6]2\text{Ca}2\text{H}_2\text{O}$
Pumpellyite:	$[(\text{Si}_2\text{O}_7)[\text{SiO}_4]_2(\text{OH})_3\text{O}]$ $(\text{Al}, \text{Fe}^{3+})_3(\text{Mg}, \text{Fe}^{2+})\text{Ca}_42\text{H}_2\text{O}$	Scapolite:	Marialite $[\text{Si}_3\text{AlO}_8]_3\text{ClNa}_4$
Garnet:	Grossularite $[\text{SiO}_4]_3\text{Al}_2\text{Ca}_3$		Meionite $[\text{SiAlO}_4]_6\text{CO}_3\text{Ca}_4$

TABLE 1

The domains defined by the margarite-häüyne-[epidote/pumpellyite], häüyne-prehnite--[epidote/pumpellyite] and häüyne-prehnite-pectolite junctions are susceptible of being described by a large number of mineral phases which are stable under the temperature and pressure conditions under consideration and close to the compositions of the natural basic rocks which are generally within the margarite-quartz-diopside triangle.

Knowing that the composition of the oxides of the Portland cement is as given in Table 2 (weight %):

CaO	SiO ₂	Fe ₂ O ₃	Al ₂ O ₃	SO ₃	MgO	Na ₂ O	K ₂ O
64.2	19.4	5.5	4.5	2.8	2.0	0.1	0.6

TABLE 2

The composition has to be enriched with alumina and silica to place it into these favourable regions. However, the composition has to be able to be prepared with a quantity of mixing water such that the density of the slurry is of the order of the

densities usually used for cementing an oil well or the like and the slurry must have rheological properties of the slurry must be compatible with pumping in such a well.

5 Six slurries with a density of 1.95 g/cm^3 and one of 1.49 g/cm^3 were thus prepared based on class G Portland cement (Black label). The formulations are shown in Table 3 below. For the liquid constituents (fluid loss control agents and antifoaming agent), the quantities are indicated in gallons per sack of Portland cement (1 gps = 3.78 litres per 94 pound cement sack, i.e., 1 gps = 0.070 litres per kilogram of Portland cement). For all of the other constituents (the solid constituents) the
10 quantities are given as the percentages by weight with respect to the weight of Portland cement (BWOC = by weight of cement).

All of the proposed formulations were optimised to obtain a slurry with a rheology compatible with pumping in a subterranean well, which necessitates adding
15 dispersing agents.

Composition #1 was a prior art oil-well cement composition intended for high temperature wells, containing 35% silica. The other compositions were compositions of the invention comprising 30% to 80% of micro-alumina and 75% to 85% of silica
20 in the form of a mixture of silica with a grain size close to the grain size of a Portland cement (average particle size close to $20 \mu\text{m}$ for silica and $14 \mu\text{m}$ for cement) and micro-silica (average particle size close to $1.2 \mu\text{m}$). For composition 7, silica and alumina are also present in the form of silica/alumina microspheres having a diameter of about $100\mu\text{m}$.

25

#	1	2	3	4	5	6	7
Micro-alumina	-	70	60	80	70	60	14
Silica	35	55	55	55	85	85	-
Micro-silica	-	30	30	30	-	-	52
Silica/alumina microspheres	-	-	-	-	-	-	44
Dispersing agent	0.3	0.8	0.8	0.8	0.8	0.8	0.03
Retarder 1	0.3	0.5	0.45	0.55	0.9	0.9	0.44
Retarder 2	0.15	0.3	0.25	0.33	0.5	0.5	0.26
Fluid loss control agent	0.6	0.6	0.6	0.6	0.6	0.6	2.5
Antifoaming agent	0.05	0.05	0.05	0.05	0.05	0.05	0.03

TABLE 3

The relative amounts of silicon, aluminium and calcium in these different formulations are as follows:

#	1	2 or 5	3 or 6	4	7
Silicon	46%	40%	43%	38%	48%
Aluminium	5%	32%	30%	37%	20%
Calcium	49%	28%	27%	25%	33%

TABLE 4

Table 5 shows the results obtained regarding the rheology of the slurry, assuming the fluid to be a Bingham fluid (plastic viscosity and yield stress T_y). The measurements were carried out under the conditions recommended by the API (American Petroleum Institute), at ambient temperature representing the nature of the surface during mixing of the cement, and at a temperature of 149°C (300°F), representing the temperature of a cement slurry circulating at the bottom of an oil well. The gel strength was measured after leaving the slurry to stand for 10 minutes following stirring for 1 minute.

The values shown in Table II indicate that the compositions were well suited to cementing an oil well or the like.

	Rheology after mixing at ambient temperature		Rheology at bottom hole circulation temperature (145°C)		
	PV cP or mPa.s	Ty lbf/100ft ² (Pa)	PV cP or mPa.s	Ty lbf/100ft ² (Pa)	Gel strength lbf/100ft ² (Pa)
1	602	13 (6.2)	196	4 (1.9)	12 (5.7)
2	53	21 (10.1)	30	15 (7.2)	36/19 (17/9)
3	58	21 (10.1)	36	13 (6.2)	33/15 (16/7)
4	48	21 (10.1)	28	15 (7.2)	35/17 (17/8)
5	55	16 (7.7)	26	11 (5.3)	22/10 (10/5)
6	63	17 (8.1)	29	12 (5.8)	24/11 (11/5)
7	257	29 (14)	109	3 (1.45)	21/6 (10/3)

TABLE 5

Table 6 below summarises the characteristic cement setting data and the compressive strengths measured after 24 hours setting. With the exception of the free water volume measurement, carried out at laboratory temperature, all of the measurements were made at 149°C (300°F). The pumpability time or thickening time corresponded to production of a consistency, measured in standardised BC units, of 100 BC; beyond this a cement slurry is no longer considered to be pumpable. The transition time corresponds to passage from a consistency of 30 BC to a consistency of 100 BC; in other words, the transition time is characteristic of the first cement setting phase.

#	1	2	3	4	5	6	7
Free water (ml/250 ml)	0.5	0	0	0	1		0
Pumpability time (hrs:min)	3:00	5:09	4:20	4:54	4:52	5:05	4:54
Transition time	0:10	0:15	0:10	0:15	0:14	0:12	0:02
Setting time for a compressive strength of							
...50 psi [0.34 MPa]	3:50	6:02	6:00	6:06	8:37		4:40
...500 psi [3.44 MPa]	4:38	6:55	6:52	7:06	9:47		5:53
Compressive strength at 24 h psi (MPa)	5028 (34.7)	2420 (16.7)	2560 (17.7)	2083 (15.7)	3283 (22.6)		2100 (14.5)

TABLE 6

The formulations tested were perfectly stable and no tendency to sedimentation was observed, as shown by the free water volumes.

After setting for 24 hours, the compressive strength of the cements of the invention was much lower than that of a class G Portland cement but the level remained correct, while in practice, the compressive strength development was far from being complete at that time. From this viewpoint, comparing tests #2 and #5 is highly significant as the two samples differed only in the type of silica used: with a more reactive micro-silica, development was more rapid but in the first hours, the compressive strength after 24 hours was substantially lower.

High temperature-high pressure cement setting

For these tests, the cement was introduced in the form of a slurry into gold tubes 5 cm long and 0.32 cm in diameter, sealed at the end by an electric arc. The tubes were then placed for 10 days in autoclaves mounted on a 200 MPa pressure line where the pressure was regulated to $\pm 1\%$ and the temperature to $\pm 1^\circ\text{C}$. The temperature was raised over 20 minutes and cooling was carried out with compressed air over 15 minutes.

A mineralogical analysis carried out by X ray diffraction produced the following results:

Following results.

Temperature	250°C						300°C						350°C					
Pressure (MPa)	30		60		100		30		60		100		30		60		100	
#	2	3	2	3	2	3	3	2	3	2	3	2	3	2	3	2	3	
Quartz	+	+	+	*	+	+	+	++	+	+	+	-	-	-	-	-	-	
Corundum	++	++	++	++	++	++	++	++	++	++	++	-	-	-	-	-	-	
Anhydrite	+	+	+	+	+	+	+	-	*	+	-	-	-	-	-	-	-	
Xonotlite	++	+	+	++	++	+	+	++	+	++	++	+	+	+	+	*	+	
Tobermorite	+	+	+	-	+	-	-	+	-	+	+	-	-	-	-	-	-	
Anorthite	-	-	-	*	-	+	*	-	+	-	-	++	++	++	++	++	++	
Epidote I	*	*	*	*	*	*	*	*	*	*	*	+	-	*	*	-	+	
Prehnite	-	*	*	-	*	*	*	*	*	-	*	*	-	*	-	-	-	
Garnet	*	*	*	*	*	*	*	-	*	*	*	*	*	*	*	*	*	
Amorphous	++	++	++	++	++	++	++	+	+	++	++	+	+	+	+	+	+	

TABLE 7

TABLE 7

The following symbols are used in Table 7 and in the other tables below reporting mineralogical analyses,:

++: represents a large quantity;

+: represents a small quantity;

*: trace;

-: absence

It should be noted that the X ray peaks corresponding to traces of epidote I, garnet and prehnite were very low in intensity or partially superimposed on those of other minerals so that identification of these compounds was uncertain. The large quantities of quartz and corundum observed for the samples at 250°C and 300°C show that the systems have not completely reacted after 10 days. In contrast, these relic minerals disappeared at 350°C. The anhydrite disappeared and the quantity of amorphous material reduced with an increase in temperature.

Formulation 3 tended to equilibrium between xonotlite and anorthite; it was close to this junction, as can be seen in Figure 1. Formulation 2 contained other minerals (Figure 1). Microprobe analysis showed the presence of prehnite ($\text{Si}_3\text{Al}_2\text{Ca}_2$), an epidote with composition $\text{Si}_{1.4}\text{Al}_{1.6}\text{Ca}$ and a garnet with composition $\text{Si}_{3.4}\text{Al}_2\text{Ca}_3$.

Prolonged high temperature-high pressure setting of cements

A cement slurry was prepared, placed in a reactor and heated for 3 days at 120°C and 20 MPa. The fourth day, the temperature was raised to 290°C over 6 hours to prevent cracking by thermal shock. The cement slurries were kept under these conditions for two months. Mineralogical analysis by X ray diffraction is shown in Table 8:

Formulation	1	2	3	7
Corundum	-	++	+	-
Quartz	-	+	*	-
Xonotlite	++	+	+	*
Anorthite	-	++	++	+
Prehnite	-	+	*	-
Epidote 2	-	*	*	-
Wairakite	-	-	-	+

TABLE 8

After two months at 290°C, the cements of formulations 2 and 3 had not completely reacted as demonstrated by the residual presence of quartz and corundum. However, this duration was sufficient to enable the cement to continue setting by encouraging anorthite crystallisation. Garnet could not be detected in any analysis. Mineralogical equilibrium of formulations 2 and 3 was reached by three minerals:

anorthite, epidote and prehnite. That of formulation 7 was reached by anorthite and wairakite according to its position in the Si/Al/Ca phase triangle.

Formulation 2 falls well within the phase triangle by favouring prehnite and epidote to the detriment of xonotlite. Formulation 3 encourages the anorthite-xonotlite equilibrium in accordance with its position on the mineralogical junction.

Transformation of cement in the presence of brine

The sample pieces obtained above underwent attack by a chlorinated brine C constituted by an aqueous solution of chlorine salts in the following proportions:
NaCl: 199.7 g/l; $MgCl_2 \cdot 6H_2O$: 1.14 g/l; $CaCl_2$: 63.0 g/l and KCl: 22.9 g/l.

The mineralogical phases found are shown in Table 9.

Formulation	#2			#3		
Conditions	Set 10 days	Set 2 months	Brine 7 days	Set 10 days	Set 2 months	Brine 7 days
Quartz	+	+	-	+	*	-
Corundum	++	++	-	++	+	-
Xonotlite	»	+	+	+	+	+
Anorthite	-	++	++	-	++	++
Epidote 1-2	*	*	*	*	*	*
Prehnite	-	+	+	*	*	*
Garnet	*	-	*	*	-	*

TABLE 9

This produces an evolution of the mineralogical repertoire of the cement over time and as a function of the external conditions (pressure, temperature, and brine).

Attack of brine on the cement set under controlled conditions for two months causes complete reaction, as shown by the disappearance of the quartz and corundum.

Formulation 3 tends towards an anorthite-xonotlite mineralogical equilibrium. Formulation 2 developed more epidote, garnet and especially prehnite phases.

Cement setting in the presence of brine.

Cement setting tests were carried out using the same protocol as that described above but partially filling the gold tube with chlorinated brine C or a sulphated brine S, a saturated aqueous solution of sodium sulphate Na_2SO_4 . Because of the small volume of slurry in each tube, this was equivalent to using brine as mixing water.

The mineralogical composition of the cements obtained from formulation #2 after 10 days in the autoclave is given in the table below:

Temperature	250°C					300°C				350°C		
Pressure (MPa)	16		30			16		30		30		
Brine	C	S	-	C	S	C	S	C	S	-	C	S
Quartz	+	*	+	++	-	-	-	*	-	-	*	-
Corundum	++	*	++	++	-	++	-	+	-	-	+	-
Anhydrite	-	+	+	-	+	-	-	-	+	-	-	+
Xonotlite	++	-	++	+	-	++	-	+	-	+	*	-
Tobermorite	+	-	+	++	-	-	-	-	-	-	-	-
Anorthite	-	-	-	-	-	+	-	++	-	++	++	-
Epidote I	-	-	*	-	-	-	-	-	-	+	-	-
Analcime	*	-	-	*	-	-	-	-	-	-	-	-
Prehnite	*	*	-	*	*	-	*	*	*	*	*	*
Garnet	-	-	*	-	-	-	*	*	*	*	*	*
Felspathoid	-	++	-	-	++	-	++	-	++	-	-	++
Amorphous	++	++	++	++	+	++	++	++	++	+	+	+

TABLE 10

Setting of slurry in the presence of brine shows an adaptation of the cement to briny media.

The sulphated brine allowed the cement to react better than the chlorinated brine.

It can be seen that the reactivity was higher in the presence of brine, corundum and quartz disappearing earlier under identical pressure and temperature conditions. The new-formed phases were of the same nature as those obtained by chemical modification due to attack on the set cement by brine. When the brine penetrates into the set cement under pressure, the cement retains its correct mineralogical repertoire but ion exchanges occur between the brine and the minerals being formed. In contrast, when the brine is present at the time the slurry sets, it modifies the chemical composition of the cement before it sets. A stable system is thus formed from the start which is an indication of good behaviour.

Brines modify the chemical composition of cement formulations.

These first qualitative results were verified for formulation #1 of the prior art and for formulations #2 and #5. For these tests, maturing of a slurry prepared with water or a brine as the mixing water were compared, as indicated in the table below. The compressive strength (in MPa) was measured after one month of setting at 287.7°C (550°F), except for the second column of results, entitled "water + attack brine C", where the samples, prepared with water as the mixing water, were placed in the setting cell for one month before placing them in the presence of brine C for one additional month.

The column entitled "Brine C 2 months" corresponds to a sample prepared with brine C as mixing water that is placed in the setting cell for one month before placing them in the presence of brine c for one further month.

The brines used were chlorinated brine C, sulphated brine S and a mixed brine M, constituted by an aqueous solution and salts in the following proportions: Na_2SO_4 : 199.7 g/l; NaCl: 1.14 g/l; $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$: 63 g/l and KCl: 22.9 g/l.

Mixing water	Water	Water + attack brine C	Brine C 1 month	Brine C 2 months	Brine S	Brine M
#1	43.8	37.3	24.4	19.6	23.7	
#5	70.8	8.7	5.2		8.8	10.1
#2	15.2	15.8	11.0	10	11.6	10.7

It can be seen that the conventional formulation set in the presence of brine used directly as the mixing water had a compressive strength of only about 60% of the compressive strength of the samples prepared with tap water and which had not suffered brine attack. Further, mineralogical analysis showed the presence of xonotlite and foshagite (attack by a chlorinated brine) or kilkoanite (attack by a sulphated brine), which are known to be unfavourable to the development of compressive strength. After a further one month of attack by brine, the compressive strength is not more than 44% of the compressive strength of samples prepared with tap water that has not been subjected to attack by brine.

In the case of the preferred formulation #2 of the invention, degradation is much lower but in particular, mineralogical analysis showed a large anorthite presence, a phase which is known to be stable: in other words, it appears that the

majority of the maturing due to attack by brines is achieved after one month while this is only accelerated with the conventional type formulation.

CLAIMS

1. A cementing composition for an oil well or the like, based on a Portland cement, silica and alumina, characterized in that the mineralogical composition of the cement matrix is included in the Si-Ca-Al triangle in one of the margarite-haüyne-[epidote/pumpellyite], haüyne-prehnite-
5 [epidote/pumpellyite] and haüyne-prehnite-pectolite composition triangles.
2. A composition according to claim 1, in which the mineralogical composition is within the Si-Ca-Al triangle approximately in the zone delimited by silica between 35% and 50%, aluminium between 20% and 38% and calcium
10 between 25% and 35%.
3. A composition according to any preceding claim, characterized in that all or a portion of the silica is added in the form of a micro-silica.
4. A composition according to any preceding claim, characterized in that all or a portion of the alumina is added in the form of micro-alumina.
- 15 5. A composition according to any preceding claim, characterized in that the mixing water comprises salts of the same type as those present in the brines to which the cement will be exposed.
6. A composition according to claim 5, characterized in that a sulphated brine is used to prepare the cement slurry.
- 20 7. A composition according to claim 5, characterized in that a chlorinated brine is used to prepare the cement slurry.

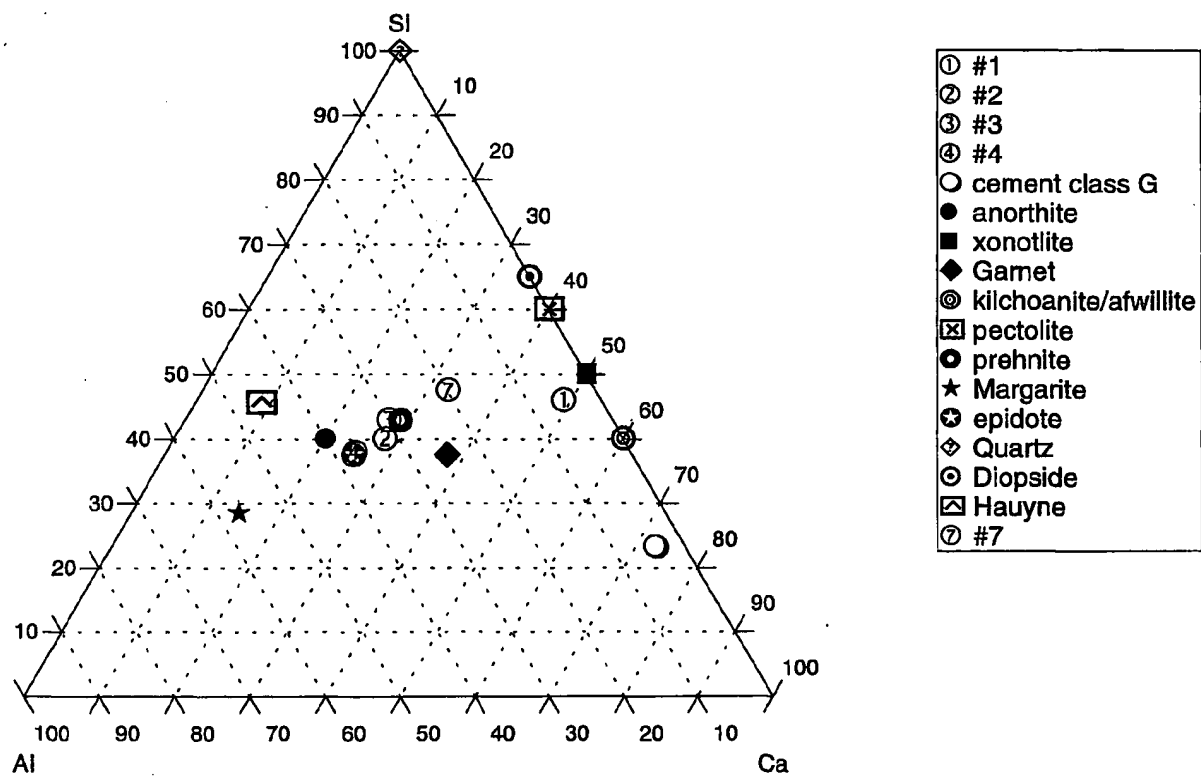
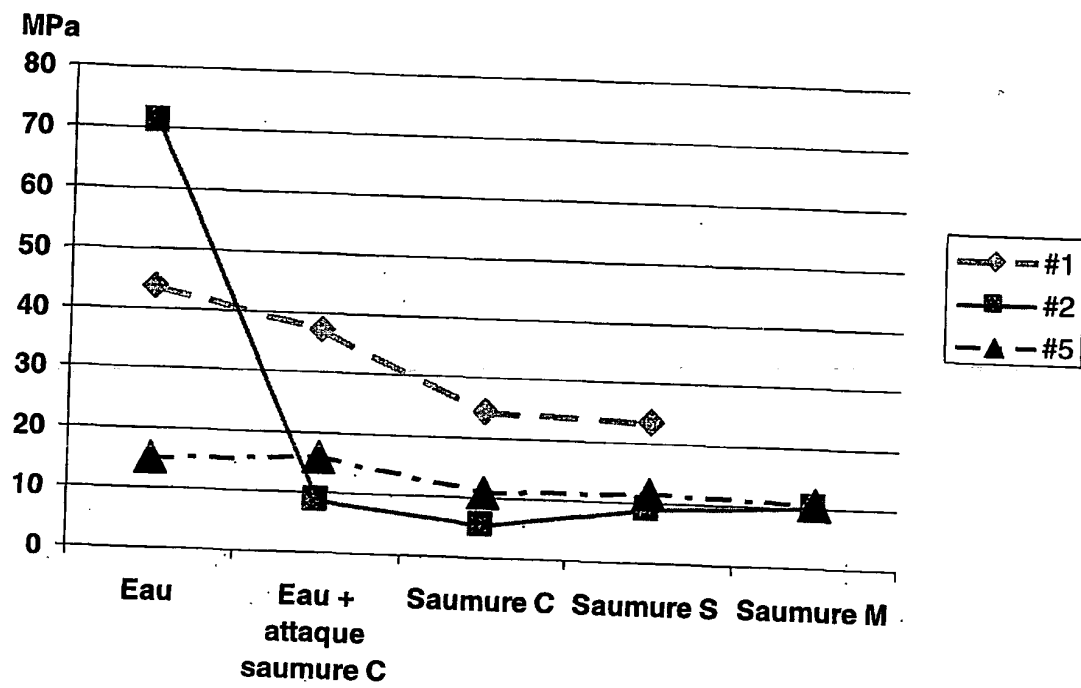


Figure 1

Figure 2



INTERNATIONAL SEARCH REPORT

National Application No

PCT/EP 01/03275

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 C04B28/04 C04B7/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C04B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal, PAJ, COMPENDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	FR 2 765 571 A (SCHLUMBERGER CIE DOWELL) 8 January 1999 (1999-01-08) cited in the application claims	1-7
A	----- CHEMICAL ABSTRACTS, vol. 108, no. 8, 22 February 1988 (1988-02-22) Columbus, Ohio, US; abstract no. 61416k, E. SAKAI, ET AL.: page 337; XP000157793 abstract & JP 62 207750 A (ID.) 12 September 1987 (1987-09-12) -----	

☐ Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

1 August 2001

Date of mailing of the international search report

09/08/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax (+31-70) 340-3016

Authorized officer

Daeleman, P

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 01/03275

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
---	---------------------	----------------------------	---------------------

FR 2765571	A	08-01-1999	AU 8540498 A	25-01-1999
			WO 9901397 A	14-01-1999
			EP 0922013 A	16-06-1999

JP 62207750	A	12-09-1987	NONE
-------------	---	------------	------